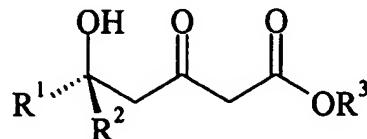


Listing of Claims

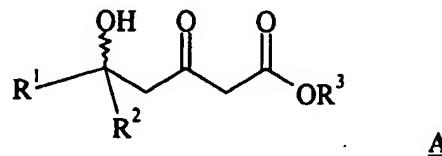
Claim 1 (currently amended): A process for preparing an optically active 5-hydroxy-3-ketoester of the formula A1 or A2



A1 or A2

or one of the tautomers thereof,

wherein R¹ and R² independently of each other represent hydrogen or a group which is selected from among C₁-C₈-alkyl, C₃-C₈-cycloalkyl, C₆-C₁₀-aryl and C₁-C₈-alkylene-C₆-C₁₀-aryl, optionally with one, two or three substituents, selected from among hydroxy, halogen, C₁-C₄-alkoxy and CF₃, where R¹ and R² do not simultaneously have the same meaning, and R³ denotes a group selected from among C₁-C₈-alkyl, C₁-C₄-Haloalkyl, C₆-C₁₀-aryl-C₁-C₈-alkylene and trihydrocarbysilyl, characterised in that a racemic mixture of a 5-hydroxy-3-ketoester of formula A



A

wherein R¹, R² and R³ are as hereinbefore defined,

is resolved into the two enantiomeric 5-hydroxy-3-ketoester A1 and A2 by preparative high performance liquid chromatography (HPLC) over a chiral carrier material, wherein the chiral carrier material is selected from the group consisting of tris(3,5-dimethylphenylcarbamate)-amylose, tris[(S)-α-methylbenzylcarbamate]-amylose, tris(3,5-dimethylphenylcarbamate)-cellulose, tris(4-methylbenzoate)-cellulose, cellulose triacetate, cellulosetribenzoate, tris(phenylcarbamate)-cellulose, tris(4-chlorophenylcarbamate)-cellulose, cellulosetricinnamate and cellulosetribenzoate.

Claim 2 (original): The process according to claim 1, wherein the two separate enantiomeric 5-hydroxy-3-ketoesters A1 and A2 are each obtained in an enantiomer excess of at least 95%.

Claim 3 (original): The process according to claim 1, wherein R¹ and R² independently of each other are selected from the group consisting of methyl, ethyl, propyl, butyl, phenyl, benzyl, phenylethyl and phenylpropyl, optionally with a substituent selected from the group consisting of hydroxy, fluorine, chlorine, bromine, methoxy, ethoxy and CF₃.

Claim 4 (original): The process according to claim 1, wherein R³ is selected from the group consisting of methyl, ethyl, propyl, butyl and benzyl.

Claim 5 (original): The process according to claim 1, wherein R¹ denotes 2-phenylethyl and R² denote propyl or R¹ denotes propyl and R² denotes 2-phenylethyl.

Claim 6 (original): The process according to claim 1, wherein R³ denotes tert.-butyl or ethyl.

Claim 7 (original): The process according to claim 5, wherein R¹ denotes 2-phenylethyl, R² denotes propyl and R³ denotes ethyl or tert.-butyl.

Claims 8-11 (cancelled)

~~8~~ **Claim 12 (currently amended):** The process according to claim ~~8~~ 1, wherein tris(3,5-dimethylphenylcarbamate)-amylose or tris(3,5-dimethylphenylcarbamate)-cellulose is used as the carrier material.

~~9~~ **13.** The process according to claim 1, wherein the preparative HPLC is used in the form of SMB (Simulated Moving Bed) chromatography.

Claims 14 and 15 (canceled)